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L1 HAS NO ANSWERS

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FILE 'HCAPLUS' ENTERED AT 11:58:30 ON 07 NOV 2009

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FILE 'REGISTRY' ENTERED AT 12:00:37 ON 07 NOV 2009

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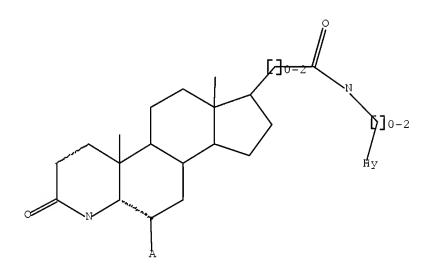
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L24 HAS NO ANSWERS L24 STR



L25 5 S L24 SSS SAM L26 130 S L24 SSS FULL

FILE 'HCAPLUS' ENTERED AT 12:11:51 ON 07 NOV 2009

L27 2 S L26

L28 1 S L27 AND (PY<2004 OR AY<2004 OR PRY<2004)

L28 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2009 ACS on STN

 ${\tt TI}$ Synthesis and uses of 4-azasteroid derivatives as selective androgen

receptor modulators (SARMs)

GI

AΒ Compds. of structural formula (I) are modulators of the androgen receptor (AR) in a tissue selective manner. They are useful as agonists of the androgen receptor in bone and/or muscle tissue while antagonizing the AR in the prostate of a male patient or in the uterus of a female patient. These compds. are therefore useful in the treatment of conditions caused by androgen deficiency or which can be ameliorated by androgen administration, including osteoporosis, osteopenia, glucocorticoid-induced osteoporosis, periodontal disease, bone fracture, bone damage following bone reconstructive surgery, sarcopenia, frailty, aging skin, male hypogonadism, postmenopausal symptoms in women, atherosclerosis, hypercholesterolemia, hyperlipidemia, obesity, aplastic anemia and other hematopoietic disorders, inflammatory arthritis and joint repair, HIV-wasting, prostate cancer, cancer cachexia, Alzheimer's disease, muscular dystrophies, premature ovarian failure, and autoimmune disease, alone or in combination with other active agents.

ACCESSION NUMBER: 2003:892539 HCAPLUS Full-text

DOCUMENT NUMBER: 139:375605

TITLE: Synthesis and uses of 4-azasteroid derivatives

as

selective androgen receptor modulators (SARMs)

INVENTOR(S): Wang, Jiabing; McVean, Carol A.

PATENT ASSIGNEE(S): Merck & Co., Inc., USA SOURCE: PCT Int. Appl., 181 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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